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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
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Peter Bernstein

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08/19/2008

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EXAMINER

O'DELL, DAVID K

ART UNIT

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1625

MAIL DATE

DELIVERY MODE

08/19/2008

PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary	Application No. 10/525,303	Applicant(s) BERNSTEIN ET AL.	
	Examiner David K. O'Dell	Art Unit 1625	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 20 May 2008.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1-4,6,7,10 and 16-28 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☒ Claim(s) 1-4,6,7,10 and 16-28 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
 2. ☐ Certified copies of the priority documents have been received in Application No. _____.
 3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- | | |
|--|---|
| 1) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892) | 4) <input type="checkbox"/> Interview Summary (PTO-413) |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948) | Paper No(s)/Mail Date. _____ |
| 3) <input type="checkbox"/> Information Disclosure Statement(s) (PTO/SB/08) | 5) <input type="checkbox"/> Notice of Informal Patent Application |
| Paper No(s)/Mail Date _____ | 6) <input type="checkbox"/> Other: _____ |

DETAILED ACTION

1. Claims 1-4, 6, 7, 10, 16-28 are pending in the current application.
2. The instant application is a 371 of PCT/SE2003/001329, filed August 26, 2003, which claims the priority of Application No. 0202567-4 filed in Sweden on August 29, 2002 and Application No. 0202986-6 filed in Sweden on October 9, 2002.

Response to Arguments

3. Applicant's arguments filed on November 9, 2007 have been fully considered but they are not fully persuasive. With respect to the 103 (a) rejection, over Harrison the examiner agrees and now wishes to make a new rejection over Stevenson and Harrison in view of Elliot and Bernstein. It is difficult for the examiner to explain why Stevenson was not previously used in the rejection and why Harrison alone was used, but the examiner agrees with the applicant that the ethers of Harrison do not make the instantly claimed compounds obvious, unless Elliot is considered. With respect to the rejection under 35 U.S.C. 112 1st paragraph for scope of enablement in regards to the compound claims the rejection is withdrawn. The rejections under 35 U.S.C. 112 1st paragraph for **lack** of enablement for treating various diseases and disorders with the compounds of the instant case is maintained for the reasons of record. The remaining claims are drawn to treating depression in its various forms and anxiety. The applicant may be correct in the interpretation of the Rosenzweig-Lipson reference, however after reviewing the state of the art anew, the previous position is further substantiated with the following citation:

“Antidepressant efficacy of the NK1 antagonist Aprepitant (MK869) (6) could be demonstrated in a placebo controlled clinical study where a dose of 300 mg (p.o., once daily) of Aprepitant was administered to patients suffering from major depressive disorder for 6 weeks. In this study, Aprepitant was well tolerated and the effectiveness of the compound as an antidepressant agent was comparable with that of the serotonin uptake inhibitor paroxetine [22, 23]. **In later studies, however, the antidepressant activity of Aprepitant could not be confirmed [24]. In summary, the existing experimental evidence suggest that substance P and the NK1 receptor are important players in the pathophysiology of central nervous diseases such as depression [18], however, the**

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partially negative results with Aprepitant are contradictory to this and additional studies will be needed to get conclusive answers on the antidepressant potential of neurokinin antagonists.” MARC GERSPACHER
“Selective and Combined Neurokinin Receptor Antagonists” *Progress in Medicinal Chemistry* **2005**, 43, 49-103

This reference serves to illustrate the very real lack of unpredictability even with clinical data. Here we have only the cell based assays, that supposedly correlate with the treatment of child abuse. The Kenakin reference was cited to show that GPCRs are actually quite complicated and not to be viewed as g-protein switches and that each conformation may be viewed as having a distinct physiological outcome. Even if we were to believe that aprepitant is useful in treating depression, that does not mean that the instantly claimed compounds are also useful as such. This receptor is a GPCR with a vast number of binding sites and conformations each of which may be associated with a distinct physiological outcome. It is specious reasoning that leads one to the conclusion that a compound which in the applicant's own words is “structurally quite different than the compounds recited in Applicants' claims” would interact with a protein in exactly the same way and lead to the same physiological outcome. The review also supports another position that while aprepitant and the applicant's compounds have no conclusive therapeutic use they are not to be viewed as worthless. The point of the article was that even though you cannot show that a compound is defined in some particular pharmacological assay, the compound should not be discarded because of the very fact that it affects the receptor means that it may be useful. This supports the examiners position that even though the compounds are not good for treating depression they still have a utility. The fact that substance P and the NK1 receptor are, in the words of Gerspacher, “important players in the pathophysiology of central nervous diseases such as depression.” has never been in dispute. The real questions are: 1) Can NK1 anatagonists treat depression? According to Gerspacher “**additional studies will be needed**

to get conclusive answers on the antidepressant potential of neurokinin antagonists.” 2)

Even if NK1 antagonists had been therapeutically useful, is it reasonable to draw conclusions from other compounds, which in the applicant's own words are “structurally quite different than the compounds recited in Applicants' claims” for the treatment of a complex disease like depression? The facts clearly show that it is not.

The applicant is correct that claims 24 and 25 were erroneously grouped with the enablement rejection. For this reason and the substantial reworking of the 103(a) rejection, the action is non-final.

Claim Rejections – 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

4. Claims 1-4, 6, 7, 24, 25 are rejected under 35 U.S.C. 103(a) as being unpatentable over Harrison et. al. U. S. patent 5,620,989 AND Stevenson, Graeme I. et. al. “4,4-Disubstituted Piperidine High-Affinity NK1 Antagonists: Structure-Activity Relationships and in Vivo Activity” *Journal of Medicinal Chemistry*, **1998**, *41*, 4623-4635 in view of Bernstein et. al. “Discovery of Novel, Orally Active Dual NK1/NK2 Antagonists” *Bioorganic and Medicinal Chemistry Letters* **2001** *11*, 2769-2773 and Elliot et. al. *Bioorganic & Medicinal Chemistry Letters* **2002**, *12*, 1755–1758. The factual inquiries set forth in *Graham v. John Deere Co.*,

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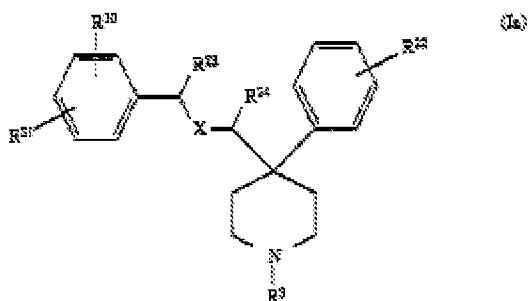
383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:

Determination of the scope and content of the prior art

(MPEP 2141.01)

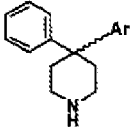
Harrison et. al. teaches NK-1 antagonists that are analogs of the compounds of the instant case that have the same utility. In particular the genus shown below:

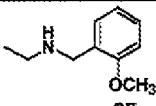
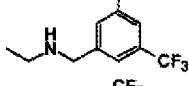
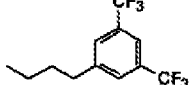
A particular sub-class of compounds according to the invention is represented by compounds of formula (Ia), and salts and prodrugs thereof:



Stevenson et. al. teach piperidinyl naphthyl amide compounds that are remarkably similar in structure and have the same utility. In particular the compounds on page 4630 Table 4:

Table 4. Alternative Linkers



Compound	Ar	hNK1 IC ₅₀ ^a	Formula	Analysis
48		> 100 ^b	C ₂₀ H ₂₇ N ₂ O	C, H, N
49		12.6 ± 8.8	C ₂₁ H ₂₂ N ₂ F ₆	C, H, N
57		63 ± 7	C ₂₂ H ₂₃ NF ₆	C, H, N*

^a Displacement of [¹²⁵I]-labeled substance P from the cloned receptor expressed in CHO cells (*n* = 3). ^b 31% and 25% @ 0.1 μM. *C₂₂H₂₃NF₆ requires 415.1734, found 415.1750.

. Elliot et. al. in his NK-1 antagonists, teaches that a variety of moieties can be used to link the 4,4-disubstituted wing carbon to an aryl group.

“Table 1 summarises the effect of modifications to the amide linker of 2. Throughout, trans-isomers have higher affinity than the corresponding cis-isomers. **A number of different linkers are tolerated, most notably the amides 2 and 15, the amine 17 and ether 23.** The relatively poor affinity of the propyl linker 24 (hNK₁ IC₅₀ 40nM) shows that a heteroatom in the chain is beneficial. This could be because the heteroatom introduces a conformational bias which favours binding, or due to participation of the heteroatom in a hydrogen bond to the receptor.” Pg. 1756-1757 Bioorganic & Medicinal Chemistry Letters 12 (2002) 1755–1758

Table 1. Linker replacements

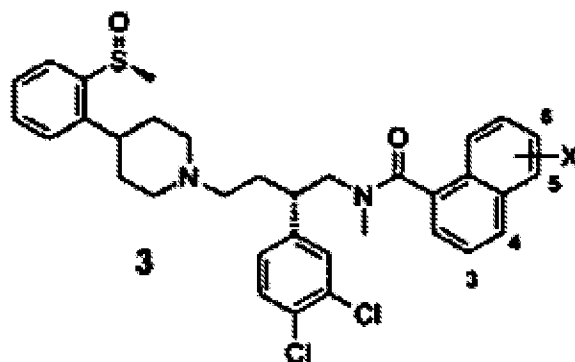
Compd	-L-	Stereochemistry	hNK ₁ IC ₅₀ (nM) ^a
11		<i>cis</i> -	150 ± 80
2		<i>trans</i> -	0.34 ± 0.10
12		<i>cis</i> -	250 ± 26
13		<i>trans</i> -	6.3 ± 2.5
14		<i>cis</i> -	85 ± 46
15		<i>trans</i> -	0.70 ± 0.44
16		<i>cis</i> -	82 ± 0
17		<i>trans</i> -	1.7 ± 0.6
18		<i>cis</i> -	140 ± 49
19		<i>trans</i> -	2.5 ± 0.6
20		<i>cis</i> -	50% @ 1000
21		<i>trans</i> -	120 ± 99
22		<i>cis</i> -	59 ± 18
23		<i>trans</i> -	4.2 ± 1.9
24		1:1 <i>cis</i> - and <i>trans</i> -	40 ± 3

^aDisplacement of [¹²⁵I]-labelled substance P from the cloned receptor expressed in CHO cells. Data are mean ± SD (n = 3).⁵

Bernstein et. al. "Discovery of Novel, Orally Active Dual NK1/NK2 Antagonists" *Bioorganic and Medicinal Chemistry Letters* **2001** *11*, 2769-2773, teaches the exact modification (down to the very precise substituents on the naphthyl ring), thus the use of naphthyl in these NK1-

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antagonists and the substitution of naphthyl for phenyl was very well known even in this very small field. Starting with the known selective antagonist SR48968, Bernstein modified the phenyl portion by screening a diverse set of compounds where the phenyl was replaced with "over 100 aryl, heteroaryl and arylalkyl groups". In the words of Bernstein the result of this study was "the most potent, dual acting compound to come out of this array was the naphthamide **2a**". Bernstein goes on to describe that "to follow-up the discovery of this naphthamide we explored the effect of substituents on the naphthalene ring."

Table 2. Exploration of varying substituents in 3-X-naphthamides

Compd	3 X=	pK_B^a NK ₁	pK_B^a NK ₂	Dose ratio ^b	
				NK ₁	NK ₂
2a	H	7.89 ± 0.08	8.18 ± 0.28	52	262
3b	NO ₂	8.16 ± 0.10	9.03 ± 0.18	50	321
3c	Br	8.15 ± 0.34	7.67 ± 0.24	43	34
3d	C≡N	8.98 ± 0.17	8.26 ± 0.10	144	74
(ZD6021)					
3e	SO ₂ CH ₃	7.43 ± 0.25	7.35 ± 0.04	22	28
3f	Cl	7.15 ± 0.12	7.10 ± 0.09	13	31
3g	OMe	7.95 ± 0.04	7.70 ± 0.06	47	77
3h	CO ₂ H	5.68 ± 0.14	6.86 ± 0.11	ND ^c	ND
3i	CH ₃	8.03 ± 0.04	7.29 ± 0.21	26	123
3j	CH ₂ CN	8.42 ± 0.24	6.99 ± 0.06	133	39
3k	Ac	7.41 ± 0.35	7.17 ± 0.13	41	156
3l	C(=CH ₂)CH ₃	7.24 ± 0.19	7.24 ± 0.29	31	75
3m	SO ₂ NH ₂	7.54 ± 0.04	7.02 ± 0.21	170	7
3n	CON(Me) ₂	5.17 ± 0.22	7.31 ± 0.33	ND	ND
3o	C≡CH	7.71 ± 0.14	7.44 ± 0.22	23	34
3p	F	7.90 ± 0.07	8.15 ± 0.23	12	52
3q	CF ₃	7.84 ± 0.07	6.45 ± 0.25	ND	ND

It is interesting that 3-cyano-naphthyl amide was the preferred substituent, as in compound 4.

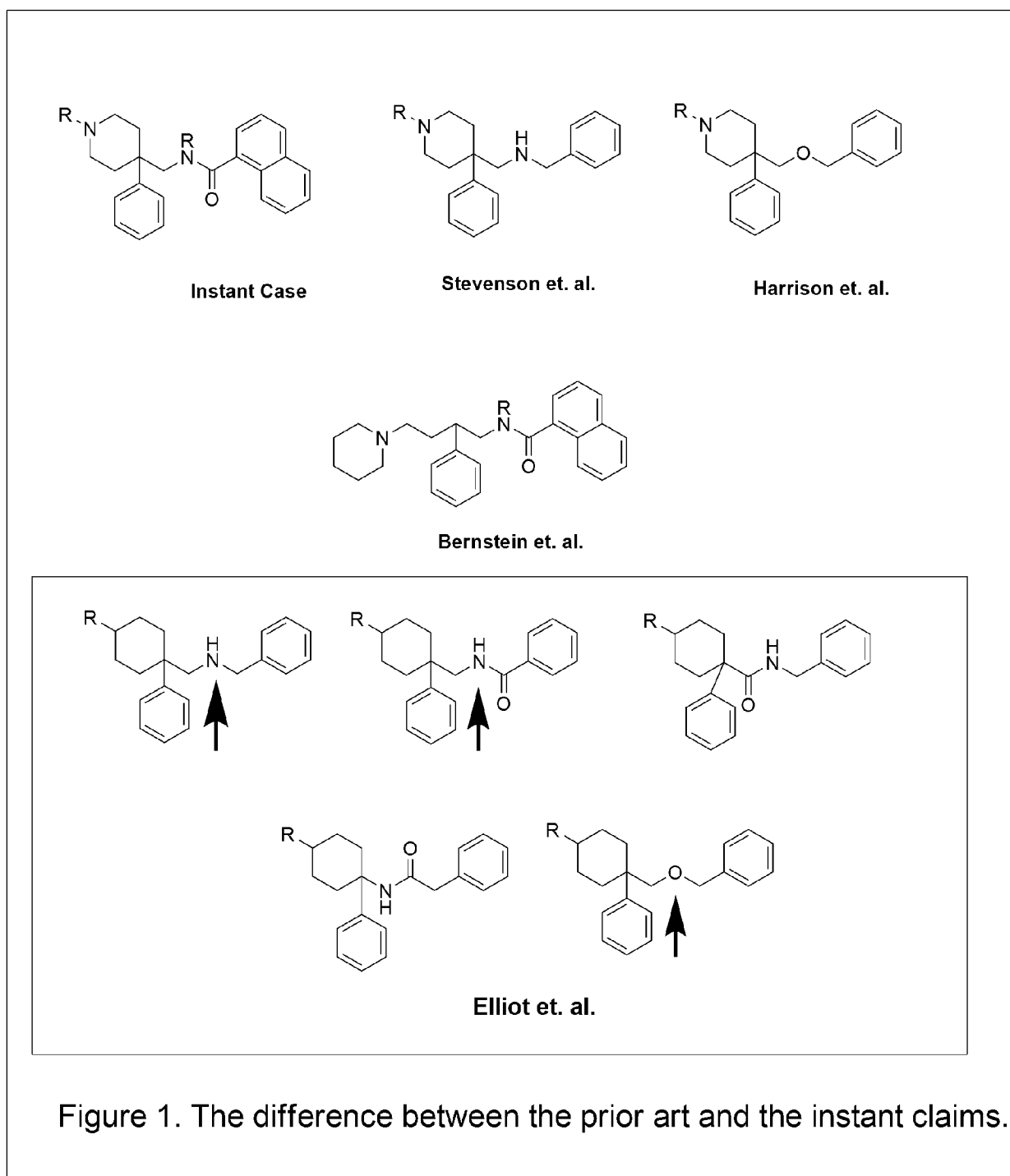
This 3-cyano naphthyl amide group is also the preferred substituent of the instant case and is

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what distinguishes these compounds from those of Stevenson et. al. and Harrison. There can be no doubt that this was the preferred substituent.

The difference between the prior art and the claims

The instant claims differ from the compounds of Stevenson et. al only in the substitution of a naphthoyl group for the benzyl group of Stevenson. These changes may also be viewed as a replacement of the O-Bn of Harrison with a naphthyl amide. This is shown below in Figure 1.



(MPEP 2141.02)

Stevenson et al. and Harrison et. al. do not expressly teach the exact compounds of the instant case.

Finding of prima facie obviousness

***Rational and Motivation
(MPEP 2142-2143)***

It would have been obvious to one of ordinary skill in the art at the time the claimed invention was made to use analogs of those of Harrison and Stevenson et. al. to produce the instant invention. Analogs differing only in the substitution of phenyl for naphthyl, are *prima facie* obvious, and require no secondary teaching when the utility is the same. The other difference the use of a different linker between the piperidine ring and the phenyl group, which are taught by Elliot as being equivalent. The experienced Ph.D. synthetic organic chemist, who would make Applicants' compounds, would be motivated to prepare these compounds on the expectation that such close analogues would have similar properties and upon the routine nature of such experimentation in the art of medicinal chemistry. It would be routine for the chemist to replace phenyl with naphthyl especially since the chemistry developed by Stevenson was in place and the synthesis only involved using an appropriate naphthyl halide (see scheme 5 of Stevenson et. al.), moreover Stevenson suggests that lipophilicity of the aryl moiety to be important since compound **49** bearing the lipophilic CF₃ group has increased potency over compound **48** (see table 4 above), thus naphthyl being slightly more lipophilic would have increased potency. Naphthyl and more specifically, the 3-cyano naphthyl group is also the preferred substituent of Bernstein et. al. who showed the preference for naphthyl over phenyl. There can be no doubt that this was the preferred substituent.

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Ex parte WESTFAHL, 136 USPQ 265 (Bd. Pat. App. & Int. 1962):

“Appellant relies upon the case of *In re Jones*, 32 CCPA 1020, 1945 C.D. 304, 579 O.G. 148, 149 F.2d 501, 65 USPQ 480, as supporting the patentability of claim 8 because in that case a naphthyl compound was held to be patentable over the corresponding phenyl compound. However, the rejection in that case was based upon the premise, held to be untenable by the court, that benzene and naphthalene are members of a homologous series. In the present case, **the examiner does not rely upon any theory of homology but has cited a reference (Richter II) teaching that naphthalene is very similar to benzene and forms a series of analogous derivatives.**”

The fact that Stevenson didn't use amides, is a non-issue since Elliot et. al. in his NK-1 antagonists replaced amino groups with amides and they were all "tolerable". It would be routine for the chemist to make the amides especially since the chemistry developed by Stevenson was in place and the synthesis only involved using an appropriate naphthyl halide or acid (see scheme 5 of Stevenson et. al.), moreover Stevenson suggests that compounds with an amino nitrogen that is too basic are less active. It goes without saying that the amino nitrogen is substantially more basic than an amide. In fact an amide is somewhat acidic. Furthermore Elliot et. al. teaches that they are in fact interchangeable.

A reference is good not only for what it teaches by direct anticipation but also for what one of ordinary skill in the art might reasonably infer from the teachings. (*In re Opprecht* 12 USPQ 2d 1235, 1236 (Fed Cir. 1989); *In re Bode* 193 USPQ 12 (CCPA) 1976). In light of the forgoing discussion, the Examiner concludes that the subject matter defined by the instant claims would have been obvious within the meaning of 35 USC 103(a). From the teachings of the references, it is apparent that one of ordinary skill in the art would have had a reasonable expectation of success in producing the claimed invention. Therefore, the invention as a whole was *prima facie* obvious to one of ordinary skill in the art at the time the invention was made, as evidenced by the references, especially in the absence of evidence to the contrary.

One of ordinary skill is also one of “ordinary creativity, not an automaton”. See *Leapfrog Enterprises Inc. v. Fisher-Price. and Mattel Inc.* UNITED STATES COURT OF APPEALS FOR THE FEDERAL CIRCUIT “An obviousness determination is not the result of a rigid formula disassociated from the consideration of the facts of a case. Indeed, the common sense of those skilled in the art demonstrates why some combinations would have been obvious where others would not. See *KSR Int’l Co. v. Teleflex Inc.*, 550 U.S. , 2007 U.S. LEXIS 4745, 2007 WL 1237837, at 12 (2007) (“The combination of familiar elements according to known methods is likely to be obvious when it does no more than yield predictable results.”).

6. Claims 10, 16-23, 26-28 are rejected under 35 U.S.C. 112, first paragraph, as based on a disclosure which is not enabling. The specification does not enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to use the invention. There are many factors to be considered when determining whether there is sufficient evidence to support a determination that a disclosure does not satisfy the enablement requirement and whether any necessary experimentation is “undue.” These factors include, but are not limited to the following:

- (A) *The breadth of the claims;*
- (B) *The nature of the invention;*
- (C) *The state of the prior art;*
- (D) *The level of one of ordinary skill;*
- (E) *The level of predictability in the art;*
- (F) *The amount of direction provided by the inventor;*
- (G) *The existence of working examples; and*
- (H) *The quantity of experimentation needed to make or use the invention*

In re Wands, 858 F.2d 731, 737, 8 USPQ2d 1400, 1404 (Fed. Cir. 1988).

Disclosure of the activity of the compounds and dosages that are critical or essential to the practice of the invention, but not included in the claims is not enabled by the disclosure. See *In re Mayhew*, 527 F.2d 1229, 188 USPQ 356 (CCPA 1976). The only information given as to what these compounds may do, at least in the pharmacological sense, is on pg. 12 of the

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disclosure “Individual IC_{50} values were reported, along with pIC_{50} . When the two IC_{50} 's obtained for a compound differed by more than 3-fold that compound was assayed one or two more times to redetermine the value. Compounds of the present invention exhibit a K_i in the range of 1 to 100 nM in the SERT assay and have an IC_{50} 's in the range 1 to 100 nM in FLIPR assay.” The applicant has given ranges of two orders of magnitude for each individual assay, without reference to a known compound that is an agonist/inhibitor and the variability in these assays make evaluation of therapeutic value difficult. In the instant case we do not know whether the compounds are partial agonists at the NK-1 receptor. It is possibly that some compounds are both SERT inhibitors and partially active at the NK-1 receptor and vice versa, or both potent inhibitors of SERT and potent antagonists at the NK-1 receptor. Applicant seems to believe these compounds are the later although no support has been provided for this assertion. Moreover, even if this dual activity was possessed by the compounds of the invention, one cannot predict *a priori* what the outcome of such complex pharmacological behavior would be in the complex diseases of claim 10. The article cited by the authors (Ryckmans, T., et al., Bioorg. Med. Chem.Lett. (2002), 12, 261). suggests that these kinds of compounds might be useful for treatment of depression and they may well be but no such evidence is provided in the instant case. The “how to use” requirement of 35 U.S.C. 112 are not met by disclosing a pharmacological activity of the claimed compounds if one skilled in the art would not be able to use the compounds effectively without undue experimentation (In re Diedrich (CCPA 1963) 318 F2d 946, 138 USPQ 128; In re Gardner et. al. (CCPA 1970) 427 F2d 786, 166 USPQ 138). The treatment of depression is not indicated by the state of the art, even with pre-clinical data (animal

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models), as stated in a recent review (Rosenzweig-Lipson et. al. *Pharmacology & Therapeutics* **2007**, *113*, 134-153) pg. 140 paragraph 3 sentence 2:

“Although the NK-1 antagonist aprepitant was not proven efficacious in Phase III depression trials (Keller et al., 2006), it is conceivable that the combination of aprepitant with an SSRI might result in rapid onset of antidepressant effects. To this end, NK-1 antagonists have been shown to potentiate the neurochemical effects of SSRIs in preclinical studies (Guiard et al., 2004). Whether this combination or other non-monoaminergic mechanisms will produce rapid onset antidepressant effects remains to be answered.”

Thus the state of the art in the area of these dual antagonists is murky at best. Even if there was a correlation of the pharmacological activity with a clinical manifestation, we have only *in-vitro* testing of these compounds and no *in-vivo* data. Without at least animal studies of *in-vivo* activity one cannot believe that these compounds will behave as therapeutics in those suffering from depression. Moreover, even if these compounds were evaluated simply as NK-1 antagonists, a recent review article (McLean, S. *Current Pharmaceutical Design* **2005**, *11*, 1529, pg. 1542 paragraph 3) states, that:

In summary, clinical studies with three different compounds demonstrate antidepressant efficacy in both mildly depressed as well as melancholic patients. Furthermore, the favorable side effect profile of the agents suggests a viable therapy particularly for people experiencing significant sexual side effects with currently available antidepressants. **This has to be balanced against a number of trials in which NK1 receptor antagonists failed to show activity. In addition to the previously mentioned negative trials, NKP608 another NK1 receptor antagonist was reported on the Novartis web site to have been terminated from further development for depression although it is unclear whether this was due to side effects or lack of efficacy. To date there are three positive trials in depression, one positive trial in panic, several failed trials and at least 2 negative studies.**

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It seems very unlikely that one skilled in the art would know what to do with these compounds. and the data given here do not correlate with the treatment given the mechanism that applicant alleges and the current knowledge in the art.

As per MPEP:

CORRELATION: IN VITRO /IN VIVO

The issue of “correlation” is related to the issue of the presence or absence of working examples. “Correlation” as used herein refers to the relationship between in vitro or in vivo animal model assays and a disclosed or a claimed method of use. An in vitro or in vivo animal model example in the specification, in effect, constitutes a “working example” if that example “correlates” with a disclosed or claimed method invention. **If there is no correlation, then the examples do not constitute “working examples.” In this regard, the issue of “correlation” is also dependent on the state of the prior art. In other words, if the art is such that a particular model is recognized as correlating to a specific condition, then it should be accepted as correlating unless the examiner has evidence that the model does not correlate. Even with such evidence, the examiner must weigh the evidence for and against correlation and decide whether one skilled in the art would accept the model as reasonably correlating to the condition.** In re Brana, 51 F.3d 1560, 1566, 34 USPQ2d 1436, 1441 (Fed. Cir. 1995) (reversing the PTO decision based on finding that in vitro data did not support in vivo applications). Since the initial burden is on the examiner to give reasons for the lack of enablement, the examiner must also give reasons for a conclusion of lack of correlation for an in vitro or in vivo animal model example. A rigorous or an invariable exact correlation is not required, as stated in Cross v. Iizuka, 753 F.2d 1040, 1050, 224 USPQ 739, 747 (Fed. Cir. 1985): [B]ased upon the relevant evidence as a whole, there is a reasonable correlation between the disclosed in vitro utility and an in vivo activity, and therefore a rigorous correlation is not necessary where the disclosure of pharmacological activity is reasonable based upon the probative evidence. (Citations omitted.)

After reviewing the state of the art anew, the previous position is further substantiated with the following citation:

“Antidepressant efficacy of the NK1 antagonist Aprepitant (MK869) (6) could be demonstrated in a placebo controlled clinical study where a dose of 300 mg (p.o., once daily) of Aprepitant was administered to patients suffering from major depressive disorder for 6 weeks. In this study, Aprepitant was well tolerated and the effectiveness of the compound as an antidepressant agent was comparable with that of the serotonin uptake inhibitor paroxetine [22, 23]. **In later studies, however, the antidepressant activity of Aprepitant could not be confirmed [24].** In summary, the existing experimental evidence suggest that substance P and the NK1 receptor are important players in the pathophysiology of central nervous diseases such as depression [18], however, the partially negative results with Aprepitant are contradictory to this and additional studies will be needed to get conclusive answers on the antidepressant

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potential of neurokinin antagonists.” MARC GERSPACHER “Selective and Combined Neurokinin Receptor Antagonists” *Progress in Medicinal Chemistry* **2005**, 43, 49-103

This receptor is a GPCR with a vast number of binding sites and conformations each of which may be associated with a distinct physiological outcome. One reviewer has summarized the situation this way (Terry Kenakin and Ongun Onaran “The ligand paradox between affinity and efficacy: can you be there and not make a difference?” *TRENDS in Pharmacological Sciences* **2002**, 23, 275-280):

“A probabilistic model of protein conformation can be used to quantify the probability of various receptor conformations and the effect of ligand binding on those conformations. The basic idea behind the probabilistic model is that the function of a receptor protein is not assigned to particular conformations of the receptor. Instead, the function arises as a result of ligand-induced perturbation of the distribution of conformational states over the conformational space of the receptor.....**The foregoing discussion leads to the general conclusion that if a ligand binds to the receptor, it most probably will produce a bias in the conformations of the receptor ensemble** [i.e. it will change the receptor by its presence (Fig. 3)]. Therefore, this suggests that all ligands with macro-affinity should be extensively studied for pharmacological activities other than simple G-protein activation because various physiological activities have been defined that are mediated by conformations not necessarily related only to G-protein activation..... Ligand activities that are not related to a standard G-protein-mediated physiological response **might have therapeutic utility**.

Here we have exactly this situation, namely a ligand with affinity, but no known function, which as Kenakin et. al. concluded “...the discovery of macro-affinity of a ligand for a receptor should be considered only a starting point for the optimal exploitation of a drug for therapeutic utility.”

The factors outlined in *In re Wands* mentioned above apply here, and in particular As per the MPEP 2164.01 (a): “A conclusion of lack of enablement means that, based on the evidence regarding each of the above factors, the specification, at the time the application was filed, would not have taught one skilled in the art how to use”....”the claimed invention without undue experimentation. *In re Wright* 999 F.2d 1557,1562, 27 USPQ2d 1510, 1513 (Fed. Cir. 1993).”

It is very clear that one could not make/use this very broad invention that has no working examples in this unpredictable art without undue experimentation.

Double Patenting

The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the “right to exclude” granted by a patent and to prevent possible harassment by multiple assignees. A nonstatutory obviousness-type double patenting rejection is appropriate where the conflicting claims are not identical, but at least one examined application claim is not patentably distinct from the reference claim(s) because the examined application claim is either anticipated by, or would have been obvious over, the reference claim(s). See, e.g., *In re Berg*, 140 F.3d 1428, 46 USPQ2d 1226 (Fed. Cir. 1998); *In re Goodman*, 11 F.3d 1046, 29 USPQ2d 2010 (Fed. Cir. 1993); *In re Longi*, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); *In re Van Ornum*, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); *In re Vogel*, 422 F.2d 438, 164 USPQ 619 (CCPA 1970); and *In re Thorington*, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CFR 1.321(c) or 1.321(d) may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent either is shown to be commonly owned with this application, or claims an invention made as a result of activities undertaken within the scope of a joint research agreement.

Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CFR 3.73(b).

7. Claims 1-4, 6, 7, 10, 16-28 are provisionally rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 1-13 of copending Application No. 10/539,140 in view of Elliot et. al. Bioorganic & Medicinal Chemistry Letters 12 (2002) 1755–1758.

This is a provisional obviousness-type double patenting rejection. The instant claims differ from those of the ‘140 application by the identity of the moiety linking the naphthyl ring to the piperidine. This change is taught by Elliot et. al. in his NK-1 antagonists.

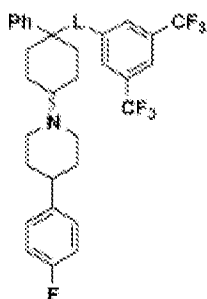
“Table 1 summarises the effect of modifications to the amide linker of 2. Throughout, trans-isomers have higher affinity than the corresponding cis-isomers. **A number of different linkers are tolerated, most notably the amides 2 and 15, the amine 17 and ether 23.** The relatively poor affinity of the propyl linker 24 (hNK₁ IC₅₀ 40nM) shows that a heteroatom in the chain is beneficial. This could be because the heteroatom introduces a conformational bias which favours binding, or

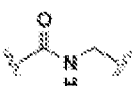
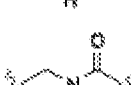
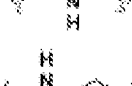
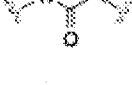
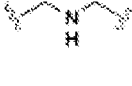
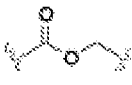
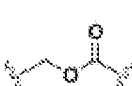
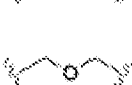


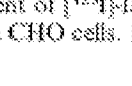




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due to participation of the heteroatom in a hydrogen

bond to the receptor." Pg. 1756-1757 Bioorganic & Medicinal Chemistry Letters 12 (2002) 1755-1758

Table 1. Linker replacements



Compd	-L-	Stereochemistry	hNK ₁ IC ₅₀ (nM) ^a
11		<i>cis</i> -	150 ± 80
2		<i>trans</i> -	0.34 ± 0.10
12		<i>cis</i> -	250 ± 26
13		<i>trans</i> -	6.3 ± 2.5
14		<i>cis</i> -	85 ± 46
15		<i>trans</i> -	0.70 ± 0.44
16		<i>cis</i> -	82 ± 0
17		<i>trans</i> -	1.7 ± 0.6
18		<i>cis</i> -	140 ± 49
19		<i>trans</i> -	2.5 ± 0.6
20		<i>cis</i> -	50% @ 1000
21		<i>trans</i> -	120 ± 99
22		<i>cis</i> -	59 ± 18
23		<i>trans</i> -	4.2 ± 1.9
24		1:1 <i>cis</i> - and <i>trans</i> -	40 ± 3

^aDisplacement of [¹²⁵I]-labelled substance P from the cloned receptor expressed in CHO cells. Data are mean ± SD (n = 3).⁵

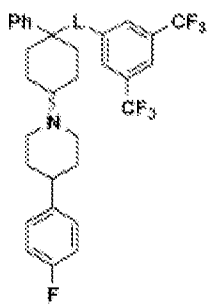
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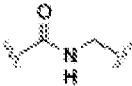
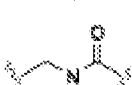
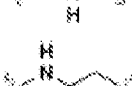


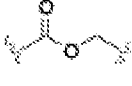
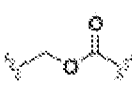
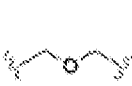

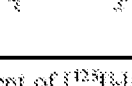
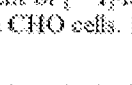
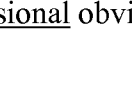



11. Claims 1-4, 6, 7, 10, 16-28 are provisionally rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 1-12 of copending Application No. 10/527,280. The claims are coextensive in scope. in view of Elliot et. al. Bioorganic & Medicinal Chemistry Letters 12 (2002) 1755–1758.

This is a provisional obviousness-type double patenting rejection. The instant claims differ from those of the '280 application by the identity of the moiety linking the naphthyl ring to the piperidine. This change is taught by Elliot et. al. in his NK-1 antagonists.

“Table 1 summarises the effect of modifications to the amide linker of 2. Throughout, trans-isomers have higher affinity than the corresponding cis-isomers. **A number of different linkers are tolerated, most notably the amides 2 and 15, the amine 17 and ether 23.** The relatively poor affinity of the propyl linker 24 (hNK₁ IC₅₀ 40nM) shows that a heteroatom in the chain is beneficial. This could be because the heteroatom introduces a conformational bias which favours binding, or due to participation of the heteroatom in a hydrogen bond to the receptor.” Pg. 1756-1757 Bioorganic & Medicinal Chemistry Letters 12 (2002) 1755–1758

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^aDisplacement of [¹²⁵I]-labelled substance P from the cloned receptor expressed in CHO cells. Data are mean ± SD (n = 3).⁵

This is a provisional obviousness-type double patenting rejection.

Conclusion

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9. No claims are allowed. Any inquiry concerning this communication or earlier communications from the examiner should be directed to David K. O'Dell whose telephone number is (571)272-9071. The examiner can normally be reached on Mon-Fri 7:30 A.M.-5:00 P.M EST.

10. If attempts to reach the examiner by telephone are unsuccessful, the examiner's Primary examiner, Rita Desai can be reached on (571)272-0684. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

D.K.O.

/Rita J. Desai/
Primary Examiner, Art Unit 1625

